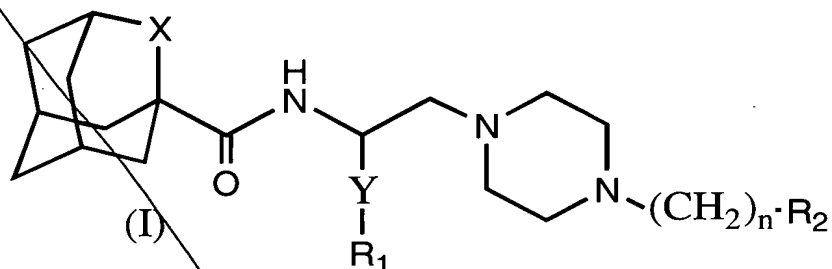


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What is Claimed:

1. A compound of the formula (I):



wherein:

X is selected from $-\text{CH}_2-$ or a chemical bond;

Y is selected from $-(\text{CH}_2)_m-$ or $-(\text{CH}_2)-\text{O}-(\text{CH}_2)-$;

m is selected from the integer 0 or 1;

n is selected from the integer 0 or 1;

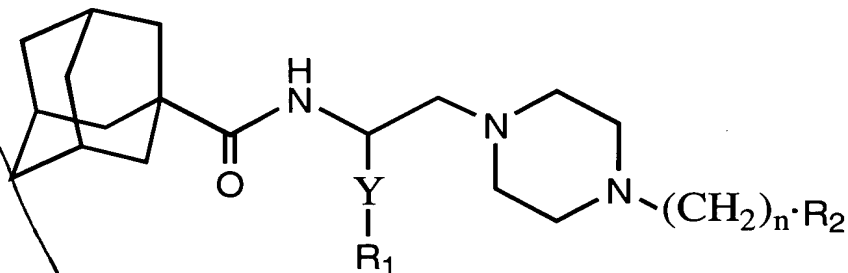
R_1 and R_2 are independently selected from the group consisting of aryl or heteroaryl of from 5 – 10 atoms optionally substituted with F, Cl, Br, I, $-\text{OH}$, $-\text{NH}_2$, $-\text{CO}_2\text{H}$, $-\text{CO}_2-\text{C}_1-\text{C}_6$ alkyl, $-\text{CN}$, $-\text{NO}_2$, C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_1-C_6 perhaloalkyl, OR_3 , or C_1-C_6 perhaloalkoxy;

R_3 is selected from the group consisting of H, C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_6-C_{10} aryl, mono or bicyclic heteroaryl, C_7-C_{14} aralkyl, and mono or bicyclic heteroaralkyl, where the aryl or heteroaryl group is optionally substituted with one to three substituents independently selected from the group consisting of F, Cl, Br, I, CN, $-\text{NH}_2$, $-\text{NO}_2$, $-\text{OH}$, alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_1-C_6 perhaloalkyl, C_1-C_6 alkoxy, and C_1-C_6 perhaloalkoxy; and the optical isomers or a pharmaceutically acceptable salt thereof.

2. A compound of Claim 1 having the formula:

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wherein:

Y is selected from $-(CH_2)_m-$ or $-(CH_2)-O-(CH_2)-$;

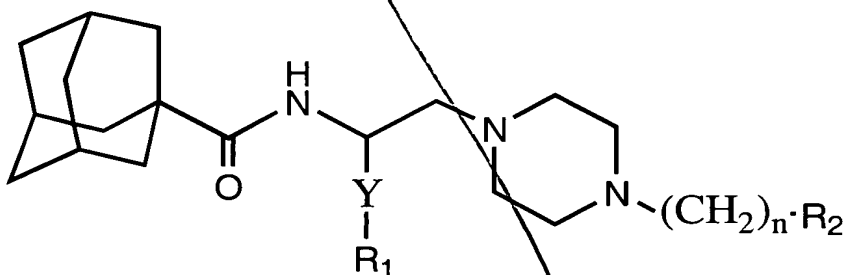
m is selected from the integer 0 or 1;

n is selected from the integer 0 or 1;

R₁ is phenyl optionally substituted with F, Cl, Br, I, -OH, -NH₂, -CO₂H, -CO₂-, C₁-C₆ alkyl, -CN, -NO₂, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ perhaloalkyl or C₁-C₆ perhaloalkoxy;

R₂ is selected from phenyl, naphthyl, piperazinyl, pyridine, thiophene, furan, imidazole, oxazole, pyrrole, pyrimidine, pyridazine, pyrazine, thiazole or oxathiazole; and the optical isomers or a pharmaceutically acceptable salt thereof.

3. A compound of Claim 1 of the formula:



wherein:

Y is selected from $-CH_2-$;

m is selected from the integer 0 or 1;

n is selected from the integer 0 or 1;

R₁ is phenyl optionally substituted with F, Cl, Br, I, -OH, -NH₂, -CO₂H, -CO₂-, C₁-C₆ alkyl, -CN, -NO₂, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ perhaloalkyl or C₁-C₆ perhaloalkoxy;

R₂ is phenyl or pyrimidinyl;

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11. A compound of Claim 1 which is (R)-N-[1-((Phenylmethoxy)methyl)-2-[4-(phenylmethyl)-1-piperazinyl]ethyl]tricyclo-[3.3.1.1^{3,7}]-decane-1-carboxamide Dihydrochloride Dihydrate

12. A compound of Claim 1 which is (R)-Adamantane-1-carboxylic acid [1-(phenylmethyl)-2-[4-(2-methoxyphenyl)-piperazinyl]ethyl]-amide Hemifumarate Hemihydrate.

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13. A compound of Claim 1 which is (R)-[1-(Phenylmethyl)-2-[4-(2-methoxyphenyl)-1-piperazinyl]-2-oxo-ethyl]-carbamic acid tert-butyl ester.

14. A compound of Claim 1 which is (R)-Adamantane-1-carboxylic acid
10 [1-(phenylmethyl)-2-[4-(2-methoxyphenyl)-piperazinyl]ethyl]-amide Hemifumarate Hemihydrate.

15. A compound of Claim 1 which is (S)-Adamantane-1-carboxylic acid
15 [1-(phenylmethyl)-2-[4-(2-methoxyphenyl)-piperazinyl]ethyl]-amide Hemifumarate Hydrate.

16. A method for treating neurodegenerative disorders comprising
administering a therapeutically effective amount of a compound of Claim 1 or a
pharmaceutical salt thereof, to a patient in need of said treatment.

17. The method of Claim 16 wherein the neurodegenerative disorder is
chronic.

18. The method of Claim 16 wherein the neurodegenerative disorder is
25 Alzheimer's Disease.

19. The method of Claim 16 wherein the neurodegenerative disorder is
Huntington's Disease.

20. The method of Claim 16 wherein the neurodegenerative disorder is
30 Parkinson's Disease.

21. The method of Claim 16 wherein the neurodegenerative disorder is
AIDS dementia.

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5 22. The method of Claim 16 wherein the neurodegenerative disorder is
retinal disease.

10 23. The method of Claim 16 wherein the neurodegenerative disorder is
epilepsy.

24. The method of Claim 16 wherein the neurodegenerative disorder is
amyotrophic lateral sclerosis.

15 25. The method of Claim 16 wherein the neurodegenerative disorder is
acute.

26. The method of claim 25 wherein the neurodegenerative disorder is
stroke.

20 27. The method of claim 26 wherein stroke is acute thromboembolic
stroke.

28. The method of claim 26 wherein stroke is focal ischemia.

29. The method of claim 26 wherein stroke is global ischemia.

30. The method of claim 26 wherein stroke is transient ischemic attack.

30 31. The method of Claim 16 wherein the neurodegenerative disorder is
ischemia resulting from a surgical technique involving prolonged halt of blood flow
to the brain.

32. The method of claim 16 wherein the neurodegenerative disorder is
head trauma.

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33. The method of claim 16 wherein the neurodegenerative disorder is spinal trauma.

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34. The method of claim 16 wherein the neurodegenerative disorder is hypoxia.

35. The method of claim 34 wherein the hypoxia is fetal hypoxia.

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36. A method of neuroprotection comprising administering a therapeutically effective amount of a compound of Claim 1, or a pharmaceutically acceptable salt thereof, to a patient in need thereof.

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37. A method of treating chronic pain comprising administering a therapeutically effective amount of a compound of Claim 1 or a pharmaceutical salt thereof, to a patient in need of said treatment.

38. The method of Claim 37 wherein the chronic pain is diabetic peripheral neuropathy.

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